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Gln Thr Glu His Ile Glu Leu  
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What is claimed is:

1. A method of depleting lipase from a sample, comprising:

contacting the sample including lipase with a probe, said probe capable of binding to the lipase to form a complex; and

separating the complex from the sample to thereby deplete the lipase from the sample

2. The method of claim 1, wherein the sample comprises a protein of interest.

3. The method of claim 1, wherein the sample comprises a polysorbate excipient.

4. The method of claim 3, wherein the polysorbate excipient is selected from polysorbate-20, polysorbate-60, polysorbate-80 or combinations thereof.

5. The method of claim 1, wherein the lipase is liver carboxylesterase-B1-like protein.

6. The method of claim 1, wherein the lipase is liver carboxylesterase-1-like protein.

7. The method of claim 1, wherein the probe is capable of being linked to a solid support.

8. The method of claim 7, wherein the solid support is agarose beads.

9. The method of claim 7, wherein the solid support is magnetic beads.

10. The method of claim 1, wherein the probe is attached to a solid support using a ligand.

11. The method of claim 10, wherein the ligand can be an indicator, biotin molecule, a modified biotin molecule, a nuclei, a sequence, an epitope tag, an electron poor molecule or an electron rich molecule.

12. The method of claim 1 further comprising recovering the lipase from the complex.

13. A method of a method of purifying a sample having a protein of interest and a lipase, comprising:

contacting the sample with a probe, said probe capable of binding to the lipase to form a complex; and

separating the complex from the sample to thereby purify the protein of interest in the sample.

14. The method of claim 13, wherein the sample comprises a polysorbate excipient.

15. The method of claim 14, wherein the polysorbate excipient is selected from polysorbate-20, polysorbate-60, polysorbate-80 or combinations thereof.

16. The method of claim 13, wherein the lipase is liver carboxylesterase-B1-like protein.

17. The method of claim 13, wherein the lipase is liver carboxylesterase-1-like protein.

18. The method of claim 13, wherein the probe is capable of being linked to a solid support.

19. The method of claim 18, wherein the solid support is agarose beads.

20. The method of claim 18, wherein the solid support is magnetic beads.

21. The method of claim 13, wherein the probe is attached to a solid support using a ligand.

22. The method of claim 21, wherein the ligand can be an indicator, biotin molecule, a modified biotin molecule, a nuclei, a sequence, an epitope tag, an electron poor molecule or an electron rich molecule.

23. The method of claim 13 further comprising recovering the lipase from the complex.

24. A method of decreasing degradation of polysorbate in a sample, comprising

contacting the sample including lipase and polysorbate with a probe, said probe capable of binding to the lipase to form a complex; and

separating the complex from the sample to thereby decreasing degradation of polysorbate in the sample.

25. The method of claim 24, wherein the sample further comprises a protein of interest.

26. The method of claim 24, wherein the polysorbate is selected from polysorbate-20, polysorbate-60, polysorbate-80 or combinations thereof.

27. The method of claim 24, wherein the lipase is liver carboxylesterase-B1-like protein.

28. The method of claim 24, wherein the lipase is liver carboxylesterase-1-like protein.

29. The method of claim 24, wherein the probe is capable of being linked to a solid support.

30. The method of claim 29, wherein the solid support is agarose beads.

31. The method of claim 29, wherein the solid support is magnetic beads.

32. The method of claim 24, wherein the probe is attached to a solid support using a ligand.

33. The method of claim 32, wherein the ligand can be an indicator, biotin molecule, a modified biotin molecule, a nuclei, a sequence, an epitope tag, an electron poor molecule or an electron rich molecule.

34. The method of claim 24 further comprising recovering the lipase from the complex.

35. A composition having a protein of interest purified from mammalian cells, surfactant and a residual amount of liver carboxylesterase B-1-like protein, wherein the residual amount of liver carboxylesterase B-1-like protein is less than about 5 ppm.

36. The composition of claim 35, wherein the surfactant is polysorbate 80.

37. The composition of claim 36, wherein the liver carboxylesterase B-1-like protein causes degradation of the polysorbate 80.

38. The composition of claim 35, wherein the composition is a parenteral formulation

39. The composition of claim 36, wherein a concentration of the polysorbate in the composition is about 0.01% w/v to about 0.2% w/v.

40. The composition of claim 35, wherein the protein of interest is selected from a group consisting of a monoclonal antibody, a polyclonal antibody, a bispecific antibody, an antibody fragment and an antibody-drug complex.

41. The composition of claim 35 further comprising one or more pharmaceutically acceptable excipients.